1. Introduction

The existence of hazardous metals in the aquatic environment has attracted extensive attention because these metals can pose serious health problems via the food chain. Chromium (Cr), resulting from various industrial processes such as electroplating, steel making, pigment manufacturing, and wood preservation, is the second most abundant inorganic groundwater contaminant. In the aquatic environment, Cr is present in the main valence states as Cr(III) and Cr(VI). Cr(VI) is a hypotoxic pollutant for humans due to its high water solubility, mobility, and carcinogenicity. 2,5 Cr(III), on the other hand, is hypotoxic, slightly soluble, has poor mobility, and is easy to be precipitated as hydroxide. 6 In accordance with the World Health Organization (WHO), the contaminant concentration for Cr(VI) in potable and surface waters should be less than 0.05 and 0.1 mg L\(^{-1}\), respectively. To remedy the Cr(VI) pollution, there is an urgent necessity to dispose Cr(VI) before pouring it into river or sea.

Various methods including adsorption, biological degradation, membrane separation, ion exchange, filtration, etc. are employed to dispose metal ions from contaminated solutions. 7-10 Among these, adsorption is regarded as an efficient way to remove metal ions from the trade effluent because this method is most economically favorable and has simple operation and high efficiency. Moreover, different adsorbents such as clays, modified activated carbon, Fe\(^{3+}\) and iron oxides, organic-inorganic composites, etc. have been applied for the disposal of Cr(VI). 11-17 However, most traditional materials are still far beyond satisfactory for the treatment of Cr(VI) from aqueous medium due to their low adsorption performance, secondary pollution, and complicated preparation methods. Therefore, the design and investigation of new materials with high performance for wastewater treatment are still necessary. In this regard, a number of researchers have tried their best to fabricate new materials with more active sites for Cr(VI) adsorption. Moreover, if Cr(VI) could be reduced to Cr(III), the toxicity might be efficiently reduced. Thus, endowing the adsorbents with reduction property may greatly enhance their Cr(VI) removal capability.

Due to its outstanding reducing property and abundant active functional groups, poly(m-phenylenediamine) (PmPD) nanoparticles seem to be one of the most prospective high-molecular polymer adsorbents. 16-18 Nitrogen atoms of amine in PmPD

The synthesis of reclaimable adsorbents with satisfactory adsorption performance and easy separation properties is necessary for environment-related applications. In this study, novel amine-functionalized magnetic Fe\(_3\)O\(_4\) (Fe\(_3\)O\(_4\)−NH\(_2\)) nanoparticles coated with poly(m-phenylenediamine) (Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs) were synthesized successfully via oxidation polymerization. The as-prepared Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs with a well-defined core−shell structure were characterized, and their extraordinary Cr(VI) removal capability was investigated. Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs exhibit high adsorption capacity (508 mg g\(^{-1}\)) and fast adsorption rate towards Cr(VI). The abundant nitrogen-containing functional groups on the surface of Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs greatly contribute to the adsorption/reduction of Cr(VI). Moreover, the intraparticle diffusion model can be used to provide a good explanation of every stage of the process. The calculated thermodynamic parameters suggest that the adsorption of Cr(VI) onto Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs is endothermic and spontaneous. Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs can be easily separated, and the regenerated adsorbents still maintain high adsorption capacity. The results imply that Fe\(_3\)O\(_4\)−NH\(_2@\)PmPDs can be regarded as a suitable material for the treatment of Cr(VI) from contaminated water.
(Fig. S1†) have a lone pair of electrons that can improve its the adsorption capacity towards Cr(vi) via complexation and reducing action. However, few studies have been carried out on their application in Cr(vi) treatment because the PmPD nanoparticles are not easy to be separated from an aqueous solution. Magnetic nanoparticles, especially Fe3O4, have attracted significant attention for their facile separation property and avoidance of precipitation with costly and vast sludge. However, easy aggregation, presence of less active sites, and difficult control in the nanoscale range for Fe3O4 nanoparticles may cause their low adsorption capacity for Cr(vi) removal (46.79 mg g\(^{-1}\)). The magnetic-core and polymer-shell structure synergistically combining the advantages of PmPD and Fe3O4 can efficiently overcome the abovementioned problems. Functional groups are expected to be coated on Fe3O4 for their ability to coordinate Cr(vi) ions. Moreover, many magnetic nanocomposites have been fabricated for Cr(vi) removal. However, their adsorption capability is far from being satisfactory. Thus, the development of core–shell magnetic nanocomposites with high adsorption capability for Cr(vi) removal is still a great challenge.

Therefore, the present study was aimed at the controllable preparation of Fe3O4 by grafting amine-functional groups from 1,6-hexanediamine via one-pot solvothermal method; this controllable preparation could impede the nanoparticles against aggregation without disturbing their desirable properties and enhance the nitrogen compounds. Then, the amine-functionalized magnetic Fe3O4 (FeO4–NH\(_{2}\)) nanoparticles decorated by PmPD (FeO4–NH\(_{2}\)@PmPDs) can further improve the nitrogen-containing functional groups for Cr(vi) adsorption and reduction. The FeO4–NH\(_{2}\)@PmPD composites can be easily separated from the contaminated media. Their adsorption behavior towards Cr(vi) was evaluated, showing exceptionally high adsorption performance and sensational recycling.

Moreover, detailed investigation of the Cr(vi) adsorbed onto FeO4–NH\(_{2}\)@PmPDs implies that Cr(vi) is reduced to Cr(II) by benzenoid amine groups on FeO4–NH\(_{2}\)@PmPDs. In conclusion, the design of FeO4–NH\(_{2}\)@PmPDs provides a novel insight into the use of core–shell nanoparticles for the Cr(vi) removal, and these FeO4–NH\(_{2}\)@PmPDs exhibit tremendous potential for Cr(vi) removal from Cr-contaminated wastewater.

2. Materials and methods

2.1 Materials

Sodium citrate (NaC\(_{6}\)H\(_{5}\)O\(_{7}\)), anhydrous sodium acetate (CH\(_{3}\)COONa), ferric chloride hexahydrate (FeCl\(_{3}\)·6H\(_{2}\)O), ethylene glycol (HOCH\(_{2}\))\(_{2}\), sodium persulfate (Na\(_{2}\)S\(_{2}\)O\(_{8}\)), 1,6-hexanedi-amine, and m-phenylenediamine (m-PD) were purchased as analytical reagents from Sinopharm Chemical Reagent Co. Ltd. and used without any purification. Storing solution including 360 mg L\(^{-1}\) Cr(vi) was provided by dissolving K\(_{2}\)Cr\(_{2}\)O\(_{7}\). Milli-Q water was used for the preparation of solutions in the experiment.

2.2 Preparation of FeO4–NH\(_{2}\) and FeO4–NH\(_{2}\)@PmPDs

FeO4–NH\(_{2}\) nanoparticles were synthesized utilizing one-pot solvothermal method via the morphology-mediated (1,6-hexanediamine) and the precursor (FeCl\(_{3}\)·6H\(_{2}\)O), which were different from those reported in some previous studies. 1,6-Hexanediamine (8 g), NaC\(_{6}\)H\(_{5}\)O\(_{7}\) (6 g), and FeCl\(_{3}\)·6H\(_{2}\)O (2 g) were dissolved in (HOCH\(_{2}\))\(_{2}\) (60 mL) under ultrasonication, and the mixture was homogenized via magnetic stirring for several hours. Then, the mixture was placed in a Teflon-lined autoclave and heated at 200°C for 10 h. The obtained sample was ultrasonically washed several times with water and ethanol and then dried in vacuum.

FeO4–NH\(_{2}\)@PmPD nanocomposites were synthesized by a modified ice-water bath method. Briefly, FeO4–NH\(_{2}\) (0.3 g) was added to the aqueous solution (300 mL) under ultrasonication for 20 min. After this, the suspension was intensely stirred at 0°C for 10 min. Then, m-phenylenediamine and Na\(_{2}\)S\(_{2}\)O\(_{8}\) were placed in the suspension while maintaining the state for 5 h. Finally, the as-prepared composites were washed with water and ethanol and dried in vacuum at 60°C. FeO4–NH\(_{2}\) nanoparticles in different mass ratios with respect to m-phenylenediamine (1 : 1, 1 : 4, and 4 : 1) were fabricated, which were named as 4 : 1FeO4–NH\(_{2}\)@PmPDs, 1 : 1FeO4–NH\(_{2}\)@PmPDs, and 1:4FeO4–NH\(_{2}\)@PmPDs, respectively.

2.2.1 Characterization. The morphologies of the as-prepared materials were characterized by high-resolution transmission electron microscopy (HRTEM, JEOL-2010, Tokyo, Japan). The functional groups of the adsorbents were investigated by Fourier-transform infrared spectroscopy (FT-IR, Nicolet 8700, Thermo Scientific Instrument, USA) at a spectral resolving power of 4 cm\(^{-1}\). Thermogravimetric analysis (TGA) was implemented using the TGA-60/60H thermal analyzer (Shimadzu, Kyoto, Japan) under a N\(_{2}\) atmosphere at a heating speed of 20°C min\(^{-1}\). Powder X-ray diffraction (XRD) patterns were obtained using a (Philips X’Pert Pro Super X-ray) diffractometer with K\(_{x}\) source (\(\lambda = 1.54178\) Å). The zeta potentials of the samples were obtained as a function of pH using a Nano ZS instrument (Malvern Instrument, UK) at 25°C. The X-ray photoelectron spectroscopy (XPS) measurements were conducted using ESCALAB 250 (Thermo-VG Scientific, USA). Vibrating sample magnetometer (VSM) was employed to test the magnetism of the as-prepared nanocomposites, and the range of the magnetic field was between –30 000 and 30 000 Oe. The core–shell nanoparticles were also investigated via Raman spectroscopy (RAMANLOG 6, SPEX company, USA) at room temperature.

2.3 Batch experiments

The adsorption experiments were performed in 10 mL polycarbonate centrifuge tubes. Different volume of adsorbents (1.5 g L\(^{-1}\)) and Cr(vi) (360 mg L\(^{-1}\)) were added to the suspensions with the desired concentrations of the solid/solution = 0.15 g L\(^{-1}\), [Cr(vi)] = 60 mg L\(^{-1}\). The suspensions were also mixed with NaCl (0.001, 0.01 or 0.1 mol L\(^{-1}\)) for ionic strength investigation. Moreover, the desired pH of the suspensions was adjusted using negligible amounts of 0.1 M HCl or NaOH. For isotherm and kinetics experiments, pH was controlled at 2.0 and the temperature was set at 313 K. To achieve the adsorption equilibrium, the suspensions were shaken for at least 12 h, and
then, a magnet was used to separate the solid from the solid-liquid mixed phase. Finally, the concentration of Cr(VI) was detected via the spectrophotometric method ($\lambda_{\text{max}} = 540$ nm) using diphenylcarbazide as the chromogenic reagent.

Recycling of $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs was tested using a 0.5 M NaOH solution as the desorbing agent at room temperature. The adsorbents were added to the NaOH solution under ultrasonication (1 h), intensely stirred (10 h), obtained via magnetic separation, and washed several times with water. The recycled adsorbents can be extracted and used for the disposal of Cr(VI) again.

The concentration of Cr(VI) adsorbed on $\text{Fe}_3\text{O}_4$–$\text{NH}_2$ or $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs was determined via the differences between the original concentration ($C_0$, mg L$^{-1}$) and final concentration ($C_e$, mg L$^{-1}$). Accordingly, the removal% representing the percentage of Cr(VI) removal as compared to the initial concentration of Cr(VI), distribution coefficient ($K_d$), and the adsorption capacity ($C_a$, mg g$^{-1}$) were expressed as follows:  

$$\text{Removal} \% = \frac{C_0 - C_e}{C_0} \times 100\% \quad (1)$$  

$$K_d = \frac{C_0 - C_e}{C_e} \times \frac{V}{m} \quad (2)$$  

$$C_a = \frac{C_0 - C_e}{m} \times \frac{V}{m} \quad (3)$$

where $V$ represents the suspension volume, $m$ is the mass of the adsorbents. Laboratory data were the mean of triplicate measurements with errors less than 5%.

3. Results and discussion

3.1 Characterization of $\text{Fe}_3\text{O}_4$–$\text{NH}_2$ and $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs

The TEM images of the as-prepared $\text{Fe}_3\text{O}_4$–$\text{NH}_2$ and $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPD nanoparticles are shown in Fig. 1. The $\text{Fe}_3\text{O}_4$–$\text{NH}_2$ nanoparticles have a mean diameter of ~50 nm and a rough surface. Moreover, they are transparent materials due to the presence of a specific ligand ($-\text{NH}_2$) on the surface of nanoparticles. The TEM image clearly shows that the core-shell $\text{Fe}_3\text{O}_4$–$\text{NH}_2$ nanoparticles are coated with PmPD. The average thicknesses of PmPD on the surface of 4 : 1Fe$_3$O$_4$–NH$_2$@PmPDs, 1 : 1Fe$_3$O$_4$–NH$_2$@PmPDs, and 1 : 4Fe$_3$O$_4$–NH$_2$@PmPDs are approximately 10, 30, and 45 nm, respectively. An HRTEM analysis indicates that the lattice spacing of 0.484 nm is well coincident with the (111) lattice plane of Fe$_3$O$_4$.

The XRD peaks of Fe$_3$O$_4$–NH$_2$ and Fe$_3$O$_4$–NH$_2$@PmPDs (Fig. 2A) at 18.35, 30.14, 35.52, 43.12, 53.42, 57.06, and 62.68° are specified to the indices (111), (220), (311), (400), (422), (511), and (440) of Fe$_3$O$_4$ by MDI jade 6.0, indicating the magnetite phase with a face-centered cubic structure. After polymerization, the characteristic diffraction peaks are well preserved; this suggests that the structure of Fe$_3$O$_4$ is not affected by the oxidation polymerization of PmPD.

To verify the chemical components and functional groups in Fe$_3$O$_4$–NH$_2$ and Fe$_3$O$_4$–NH$_2$@PmPDs, FT-IR spectroscopy was performed. For Fe$_3$O$_4$–NH$_2$ nanoparticles, as shown in Fig. 2B, the peaks are attributed to the Fe–O lattice model of Fe$_3$O$_4$ (582 cm$^{-1}$) and N–H stretching vibration (874, 1626, and 3436 cm$^{-1}$). The results imply that the amino groups were successfully grafted on the surface of Fe$_3$O$_4$. Moreover, new peaks appear in the FT-IR spectra of Fe$_3$O$_4$–NH$_2$@PmPDs, which are ascribed to the C–N stretching vibration (1270 cm$^{-1}$), quinoid ring stretching (1620 cm$^{-1}$), and benzenoid ring stretching (1500 cm$^{-1}$), confirming the coated PmPD on Fe$_3$O$_4$–NH$_2$ nanoparticles.

Fig. 1 Representative TEM, HRTEM, and EDS images. Fe$_3$O$_4$–NH$_2$ (A and E), 4 : 1Fe$_3$O$_4$–NH$_2$@PmPDs (B and F), 1 : 1Fe$_3$O$_4$–NH$_2$@PmPDs (C and G), and 1 : 4Fe$_3$O$_4$–NH$_2$@PmPDs (D and H).
nitrogen-containing functional groups on Fe₃O₄–NH₂@PmPDs can greatly contribute to Cr(VI) adsorption. After Cr(VI) adsorption (Fe₃O₄–NH₂@PmPDs-Cr), the peaks at 874 and 1500 cm⁻¹ of Fe₃O₄–NH₂@PmPDs-Cr become weaker; this suggests that Cr(VI) interacts with the nitrogen-containing functional groups. The relative intensity of quinoid imine (1620 cm⁻¹) increases due to the oxidation of benzenoid amine to quinoid imine during the reduction of Cr(VI).

The magnetization hysteresis loops were employed to investigate the saturation magnetization of the adsorbents at room temperature (Fig. 2C). The saturation magnetizations of Fe₃O₄–NH₂: 4 : 1 Fe₃O₄–NH₂@PmPDs, 1 : 1 Fe₃O₄–NH₂@PmPDs, and 1 : 4 Fe₃O₄–NH₂@PmPDs are 80.02, 66.60, 54.46, and 33.65 emu, respectively, implying a satisfactory magnetic property of the adsorbents. Therefore, the Fe₃O₄–NH₂@PmPD composites can be easily separated from the contaminated media.

The PmPD-coated Fe₃O₄–NH₂ nanoparticles were further confirmed via TGA under a N₂ atmosphere. The curve of the Fe₃O₄–NH₂ nanoparticles shows the slight weight loss (1.1%) in the range from 50 to 150 °C due to the removal of moisture and ethanol in the first step (Fig. 2D); the second weight loss (2.5%) in the range from 150 to 320 °C is due to the phase transformation of Fe₃O₄; the third weight loss (9%) from 320 to 750 °C due to the decomposition of organic moieties indicates that the surface of Fe₃O₄ is covered with abundant amino groups. For 1 : 1 Fe₃O₄–NH₂@PmPDs, the TGA presents a weight loss (9.3%) due to the removal of moisture below 150 °C and a rapid weight loss (43.4%) from 320 to 800 °C due to the decomposition of PmPD. After heating to 320 °C, the weight of the residues for Fe₃O₄–NH₂ is greater than that for 1 : 1 Fe₃O₄–NH₂@PmPDs; this confirms that PmPD is successfully grafted on the surface of Fe₃O₄–NH₂.

Raman spectra of the synthesized Fe₃O₄–NH₂ and PmPD are presented in Fig. 2E. The characteristic peaks of 331, 524, and 670 cm⁻¹ confirm that Fe₃O₄ is composed of the magnetite phase. The peaks in the range from 1300 to 1700 cm⁻¹ for Fe₃O₄–NH₂ can be attributed to the in-plane bond-stretching of sp² hybridized carbon atoms, the presence of defects, and the specific ligand (–NH₂) on the surface of Fe₃O₄–NH₂. The characteristic peaks at ~1573 and ~1352 cm⁻¹ for PmPDs are attributed to the benzenoid and quinoid bands, respectively. As observed from Fig. 2F, the characteristic peaks of Fe₃O₄ decrease or completely disappear (especially 670 cm⁻¹) in the spectrum of 1 : 1 Fe₃O₄–NH₂@PmPDs; this indicates that Fe₃O₄–NH₂ is surrounded by PmPDs. This phenomenon is similar to the change in the other core/shell nanostructures.

### 3.2 Effects of mass ratios and adsorbent contents on the Cr(VI) removal

Cr(vi) adsorbed on Fe₃O₄–NH₂@PmPDs with different mass ratios (Fe₃O₄–NH₂:PmPD) is shown in Fig. 3A. The adsorption capacity of Fe₃O₄–NH₂@PmPDs is much higher than that of Fe₃O₄–NH₂. With the increase of the PmPD loading ratio, the adsorption capacity of Fe₃O₄–NH₂@PmPDs increases. The more the addition of PmPD, the more the adsorption capacity for Cr(vi); however, the addition of PmPD does not favor the magnetic property. It should be noted that the saturation magnetization of adsorbents gradually decreases with the
coating of PmPD; thus, the mass of PmPD should be reasonably controlled. Based on these reasons, 1 : 1Fe3O4-NH2@PmPDs should be a suitable adsorbent. To further investigate the performance of the adsorbent, 1 : 1Fe3O4-NH2@PmPDs were chosen as adsorbents in the following experiments.

For a fixed initial concentration of an adsorbate, the adsorbent content is a vital factor that determines the adsorption performance of the adsorbent. Fig. 3B depicts the dependence of Cr(Ⅵ) adsorption on 1 : 1Fe3O4-NH2@PmPDs as a function of adsorbent contents ranging from 1.25 to 150 mg L⁻¹ with other variables kept constant. A meaningful result can be found that the adsorption percentage quickly increases with the increasing adsorbent contents ranging from 1.25 to 150 mg L⁻¹. When pH > 4.5, as the HCrO4⁻/Cr2O7²⁻ equilibrium is sensitive to the pH of the solution, as shown in Fig. 4B. At 2.0 < pH < 6.4, the main existing form is HCrO4⁻ (nearly 100%). When pH > 4.5, as the HCrO4⁻ species reduces, the CrO4²⁻ species increases and goes up to nearly 100% at pH 9.0. The other species maintain a small amount without changing obviously in the range of pH from 2.0 to 9.0. The zeta potential of 1 : 1Fe3O4-NH2@PmPDs, depending on the property of the particles surface, was investigated, as shown in Fig. 4C. The zeta potentials > +30 mV at pH 2.0–3.0 suggest the excellent stability of 1 : 1Fe3O4-NH2@PmPDs. The zero point of zero charge (pHpzc) value of 1 : 1Fe3O4-NH2@PmPDs is ~6.4. When pH > 6.4, Cr(Ⅵ) ions can be easily adsorbed on the surface of 1 : 1Fe3O4-NH2@PmPDs due to the negatively charged Cr(Ⅵ) ions that are deemed to show a strong attraction for the positively charged 1 : 1Fe3O4-NH2@PmPDs surface. Moreover, the complexation of Cr(Ⅵ) by amine groups accompanied by the reduction of Cr2O7²⁻ to Cr3⁺ also contributes to the Cr(Ⅵ) removal. When pH > 6.4, Cr2O7²⁻ would be repelled from the negatively charged surface, and the poor adsorption performance is caused by the lack of electrostatic affinity and the less amount of Cr2O7²⁻ reduced by benzenoid amine. Therefore, Cr(Ⅵ) removal could reach the maximum value at low pH in this study. It can be seen from Fig. 4D that the solution pH increases after Cr(Ⅵ) is adsorbed on 1 : 1Fe3O4-NH2@PmPDs. It is mainly due to the consumption of H⁺ during the protonation of amine groups (-NH2) to -NH⁺ form for the complexation of Cr(Ⅵ). Moreover, the reduction of Cr(Ⅵ) by benzenoid amine occurs along with the consumption of H⁺ or generation of OH⁻ in the reduction phase, and the reaction equations can be shown as follows:\(^{17}\)

\[
\text{Cr}_2\text{O}_7^{2-} + 14\text{H}^+ + 6\text{e}^- \rightarrow 2\text{Cr}^{3+} + 7\text{H}_2\text{O} \tag{4}
\]

\[
\text{Cr}_2\text{O}_7^{2-} + 7\text{H}_2\text{O} + 6\text{e}^- \rightarrow 2\text{Cr(OH)}_3^{+} + 8\text{OH}^- \tag{5}
\]

3.3 Effect of pH and ionic strength

The as-prepared 1 : 1Fe3O4-NH2@PmPDs were employed to study the Cr(Ⅵ) removal at various pH values. It can be seen in Fig. 4A that as the pH increases from 2.0 to 9.0, the adsorption capacity of Cr(Ⅵ) on 1 : 1Fe3O4-NH2@PmPDs decreases from approximately 380 to 40 mg g⁻¹ at C₀ (60 mg L⁻¹), and its adsorption versus pH trend is consistent with previously reported research.\(^2\) The adsorption process for Cr(Ⅵ) can be explained by the physicochemical properties of 1 : 1Fe3O4-NH2@PmPDs and the speciation distribution of Cr(Ⅵ) in water solutions in a specific pH range. The existing state of Cr(Ⅵ) is sensitive to the pH of the solution, as shown in Fig. 4B. At 2.0 < pH < 4.5, the main existing form is HCrO4⁻ (nearly 100%). The outer-sphere complexation, including electrostatic interaction, is sensitive to ionic strength, but the inner-sphere complexation is insensitive to ionic strength due to the
formation of strong chemical bonds between the adsorbate and functional groups. The weak interaction of foreign ions for Cr(VI) adsorption indicates that the adsorption process accords to the inner-sphere surface complexation. In conclusion, the results suggest that the complexation of Cr(VI) by the nitrogen-containing groups and the reduction of Cr(VI) to Cr(III) via amine groups are the principal factors promoting the adsorption.

### 3.4 Adsorption isotherm

The adsorption isotherm is regarded as one of the fundamental information about the adsorption process. The adsorption isotherms of Cr(VI) onto the 1:1Fe₃O₄–NH₂@PmPDs at 298, 313, and 328 K were obtained and are displayed in Fig. 5A. Obviously, the adsorption capacity is highest at 328 K and lowest at 298 K, implying that the increase in temperature is beneficial to Cr(VI) removal via 1:1Fe₃O₄–NH₂@PmPDs. The Langmuir, Freundlich, and Temkin models were used to fit the isotherm data. These equations are represented as follows:

**Langmuir equation:**

\[
\frac{C_s}{C_e} = \frac{1}{C_{\text{max}}b} + \frac{C_s}{C_{\text{max}}}
\]  

(6)

**Freundlich equation:**

\[
\log C_s = \log K_F + \frac{1}{n} \log C_e
\]  

(7)

**Temkin equation:**

\[
C_s = B \ln K_T + B \ln C_e
\]  

(8)

where \(b\) is the constant of the isotherm equations. \(C_{\text{max}}\) (mg g⁻¹) represents the maximum adsorption capacity of Cr(VI) onto 1:1Fe₃O₄–NH₂@PmPDs. \(K_F\) and \(1/n\) reflect the adsorption performance and the adsorption intensity. \(B\) represents the heat of adsorption. \(R\) is 8.314 J (mol K)⁻¹ and \(T\) (K) represents the temperature. \(K_T\) and \(b_T\) are related to the maximum binding energy and heat of adsorption, respectively. The obtained parameters calculated from the isotherm equations are given in Table 1. The correlation coefficients \(R^2\) of the Langmuir model are higher than those of other models for three different temperature conditions, indicating that the Langmuir model can satisfactorily depict the adsorption process. The calculated maximum adsorption capacities are 675, 610, and 508 mg g⁻¹ at 328, 313, and 298 K, respectively. The maximum adsorption capacities for 4:1Fe₃O₄–NH₂@PmPDs, 1:1Fe₃O₄–NH₂@PmPDs, and 1:4Fe₃O₄–NH₂@PmPDs were also investigated (Fig. S2 and Table S1†). Especially, the maximum mass ratios between the adsorbed Cr(VI) and the N-based group are about 3.68, 3.65, and 2.79 for 4:1Fe₃O₄–NH₂@PmPDs, 1:1Fe₃O₄–NH₂@PmPDs, and 4:1Fe₃O₄–NH₂@PmPDs, respectively. Compared with those of other magnetic adsorbents, as shown in Table 2, the adsorption capacity of 1:1Fe₃O₄–NH₂@PmPDs is significantly better. The excellent adsorption performance can be attributed to the abundant nitrogen-containing groups on the Fe₃O₄ surface. The 1:1Fe₃O₄–NH₂@PmPDs can offer more active sites than the reported adsorbents, and the adsorption–reduction method is a good strategy for the removal of Cr(VI). The Freundlich model and Temkin equation can also show some available information with high credibility in mathematics. From the Freundlich model, the values of \(1/n\) in...
the range between 0 and 1 demonstrate that the adsorption process is an advantageous and chemisorption process, and the values of $K_F$ increasing in the wake of increasing temperature suggest an endothermic process. The values of $K_T$ and $b_T$ highly confirm the strong interaction between Cr(VI) and 1:1Fe$_3$O$_4$–NH$_2$@PmPDs and the adsorption process corresponding to the chemical reaction process.

The thermodynamic parameters of Gibbs free energy change ($\Delta G^0$), enthalpy change ($\Delta H^0$), and entropy change ($\Delta S^0$) were employed to explore the thermal properties of the adsorption process. The functions are represented as follows:

$$\Delta G^0 = -RT \ln K^0$$

$$\ln K^0 = \frac{\Delta S^0}{R} - \frac{\Delta H^0}{RT}$$

where $K^0$ represents the thermodynamic equilibrium constant and $\ln K^0$ is calculated via plotting $\ln K_T$ versus $C_e$ with extrapolating $C_e$ to 0. The thermodynamic parameters of Cr(Ⅵ) adsorption on 1:1Fe$_3$O$_4$–NH$_2$@PmPDs are shown in Table 3. The positive values of $\Delta H^0$ increase as the temperature increases; this implies an endothermic adsorption process of Cr(Ⅵ) adsorption on 1:1Fe$_3$O$_4$–NH$_2$@PmPDs. The values of $\Delta G^0$ also become more negative as the temperature increase; this indicates that Cr(Ⅵ) is beneficial to be adsorbed at high temperatures, and the process is spontaneous. Moreover, the positive $\Delta S^0$ suggests that the randomness increases at the solid/solution interface with Cr(VI) adsorbed on 1:1Fe$_3$O$_4$–NH$_2$@PmPDs.

### 3.5 Adsorption kinetics

The kinetics of Cr(Ⅵ) removal by 1:1Fe$_3$O$_4$–NH$_2$@PmPDs is presented in Fig. 6A. The Cr(Ⅵ) adsorption is rapid within 200 min, in which about 80% of Cr(VI) is disposed of by 1:1Fe$_3$O$_4$–NH$_2$@PmPDs; this is attributed to the abundant surface active sites for the Cr(Ⅵ) surface adsorption/reduction reaction. Then, the Cr(Ⅵ) removal considerably slow down until it reaches equilibrium at 500 min, as well as approximately 10% Cr(Ⅵ) is adsorbed due to the relatively low residual Cr(Ⅵ) interacting with the Cr-coated 1:1Fe$_3$O$_4$–NH$_2$@PmPDs. In a practical application, the optimal contact time can be selected according to the adsorption kinetics.

To further investigate the underlying kinetics of Cr(Ⅵ) adsorption onto 1:1Fe$_3$O$_4$–NH$_2$@PmPDs, the pseudo-second-order kinetic and the intraparticle diffusion model were employed to simulate these experimental kinetic data. Moreover, the parameters for kinetic equations are shown in Table 1.
These experimental kinetic data aiming at distinguishing the intraparticle diffusion through amine groups. Through amine groups.

\[ \frac{t}{C_t} = \frac{1}{k_1 C_e} + \frac{t}{C_i} \]

The pseudo-second-order kinetic model can commendably depict the entire adsorbent systems. The adsorption of Cr(VI) on 1 : 1Fe₃O₄–NH₂@PmPDs is mostly chemisorption involving the complexation of Cr(VI) and then reduction of Cr(VI) to Cr(III) through amine groups. Some of the forefathers’ research also confirm that the Cr(VI) removal can be well described by the pseudo-second-order kinetic model.

The removal of Cr(VI) on 1 : 1Fe₃O₄–NH₂@PmPDs may undergo many steps during the whole reaction process. Thus, the intraparticle diffusion model was employed to simulate these experimental kinetic data aiming at distinguishing the different stages. As observed from Fig. 6B, adsorption is not the sole step of the rate-limiting reaction because the plots of \( q_e \) versus \( t^{1/2} \) do not yield a straight line passing through the origin. The whole adsorption process can be divided into four sections as follows: (i) Cr(VI) rapidly reaches the external surface of 1 : 1Fe₃O₄–NH₂@PmPDs on account of abundant active sites for Cr(VI) instantaneous adsorption; (ii) the adsorption slows down because the rate-limiting step for Cr(VI) diffusion into the pore microstructure or the interspace of nanoparticles; (iii) the intraparticle diffusion slows down considerably due to the low residual Cr(VI) and the interaction with the consumed functional groups; and (iv) Cr(VI) adsorption reaches equilibrium in the final stage, which is excellently stable.

### 3.6 Adsorption mechanism

XPS was employed to accurately analyze the surface chemical compositions of 1 : 1Fe₃O₄–NH₂@PmPDs before and after Cr(VI) adsorption. In the survey spectra, the peaks including C 1s, O 1s, N 1s, and Cr 2p can be observed (Fig. 7A). New peaks emerge nearby the binding energy of 577.5 eV, corresponding to the photoelectron spectra of Cr 2p, suggesting Cr(VI) adsorption on the near surface of 1 : 1Fe₃O₄–NH₂@PmPDs. Moreover, the spectra of Cr 2p shows two peaks: Cr 2p₃/₂ (577.8) and Cr 2p₁/₂ (530.4).
(587.0) via the high-resolution XPS spectrum (Fig. 7B). The broad peak of Cr 2p3/2 can be divided into two peaks at the binding energies of 578.1 and 576.3 eV, corresponding to the features of Cr(VI) and Cr(III), respectively. Moreover, it indicates that part of adsorbed Cr(vi) has been reduced to Cr(III) on the surface of 1 : 1Fe3O4–NH2@PmPDs. For comparison, the mass rates of Cr(III) and Cr(VI) adsorbed onto the surface of 1 : 1Fe3O4–NH2@PmPDs are calculated to be 53.3% and 46.7%, respectively. The changes of N 1s were also investigated by the high-resolution XPS spectra (Fig. 7C and D). Before Cr(VI) adsorption, the broad peak of N 1s can be divided into two peaks at the binding energies of 399.0 eV, corresponding to the benzenoid amine groups (−N−, 12%), and 399.79 eV, corresponding to the quinoid imine groups (−NH−, 88%), indicative of the grafted PmPD on the surface of Fe3O4. In addition, the mass fractions of benzenoid amine groups and quinoid imine groups for 1 : 1Fe3O4–NH2@PmPDs are about 2.01% and 14.70%, respectively (Table S1†). New peak appears at the binding energy of 400.58 eV, corresponding to −N=− (35%) after Cr(vi) adsorption. The phenomenon indicates that the increased −N= can be produced by −NH− during the course of reduction of Cr(vi) to Cr(III). Moreover, −N=− is derived from the in situ doping of H+ and positive Cr(III) on quinoid imine. The consumption of H+ also contributes to the increase of solution pH after Cr(vi) adsorption. The results agree well with the FT-IR analysis results stating that the adsorption of Cr(vi) occurs via the nitrogen-containing functional groups, and then, Cr(vi) is reduced to Cr(III) following the oxidation of benzenoid amine to quinoid imine.

Based on the abovementioned analyses and discussion, the Cr(vi) adsorption mechanism is illustrated in Fig. 8. The Cr(vi) adsorption involves the following steps: (i) Cr(vi) species are adsorbed by the abundant surface active sites from −NH2 and PmPD via the inner-sphere surface complexation; (ii) Cr(vi) is reduced to Cr(III) by benzenoid amine groups from PmPD; and (iii) Cr(III) readily interacts with imino groups through coordination.

### 3.7 Recycle and desorption performance

The recycle performance of 1 : 1Fe3O4–NH2@PmPDs is a principal factor to assess the cost-effectiveness of the adsorbents. Desorption of the adsorbed Cr(vi) from 1 : 1Fe3O4–NH2@PmPDs was carried out with NaOH (1mol L−1) because the affinity of 1 : 1Fe3O4–NH2@PmPDs gradually weakened with the increasing pH. As observed from the Fig. 9, the adsorption capacities of five cycles are 389, 367, 343, 337, and 332 mg g−1, respectively. The results indicate that the adsorption capacity still retains a high level during the repeated adsorption and desorption operations. The recovered nitrogen-containing functional groups and the remaining imino groups of Fe3O4–NH2@PmPDs can act as the surface active sites for the adsorption of Cr(vi) in the next cycle. The inset image of the
Fig. 9 Recycling of $1: 1\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs for Cr(Ⅵ) removal using 0.01 mol L$^{-1}$ NaOH.

separation using an external magnet suggests that $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs can be easily separated from the solution.

4. Conclusions

In this study, $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs were successfully synthesized with a uniform well-defined core-shell structure for the removal of Cr(Ⅵ) from the Cr-contaminated solutions. The as-prepared $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs enable a rapid magnetic separation and exhibit abundant nitrogen-containing functional groups towards Cr(Ⅵ) adsorption/reduction. The adsorption capacity (508 mg g$^{-1}$) of $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs at 298 K suggests an efficient Cr(Ⅵ) removal than those of other magnetic adsorbents, and the thermodynamic experiments reveal that the adsorption process is spontaneous and endothermic. The whole adsorption process follows the intraparticle diffusion model, which can be divided into four reaction stages. The adsorption of Cr(Ⅵ) on the surface active sites of $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs and the reduction of Cr(Ⅵ) to Cr(Ⅲ) via benzenoid amine are suitable to explain the removal of Cr(Ⅵ) by $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs. Recycle experiments suggest that the adsorption capacity of $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs still maintains a high level for Cr(Ⅵ) removal. Therefore, $\text{Fe}_3\text{O}_4$–$\text{NH}_2$@PmPDs could be a promising potential candidate for the treatment of Cr(Ⅵ)-contaminated water.

Conflict of interest

The authors declare no competing financial interest.

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